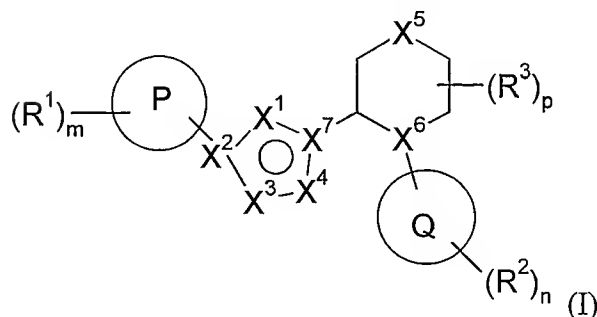


CLAIM AMENDMENTS

This listing of claims will replace all prior versions, and listings, of claims in the application.

Listing of Claims:

1. (Currently Amended) A compound according to formula I



wherein

P is selected from ~~aryl and heteroaryl~~ phenyl;

R¹ is attached to P via a carbon atom on ring P and is selected from the group consisting of ~~hydroxy~~ hydrogen, halo, nitro, C₁₋₆alkylhalo, OC₁₋₆alkylhalo, C₁₋₆alkyl, OC₁₋₆alkyl, C₁₋₆alkylOR⁵, C₀₋₆alkylcyano and C₀₋₆alkylNR⁵R⁶, C₂₋₆alkenyl, OC₂₋₆alkenyl, C₂₋₆alkynyl, OC₂₋₆alkynyl, C₀₋₆alkylC₃₋₆cycloalkyl, OC₀₋₆alkylC₃₋₆cycloalkyl, C₀₋₆alkylaryl, OC₀₋₆alkylaryl, CHO, (CO)R⁵, O(CO)R⁵, O(CO)OR⁵, O(CNR⁵)OR⁵, C₁₋₆alkylOR⁵, OC₂₋₆alkylOR⁵, C₁₋₆alkyl(CO)R⁵, OC₁₋₆alkyl(CO)R⁵, C₀₋₆alkylCO₂R⁵, OC₁₋₆alkylCO₂R⁵, C₀₋₆alkyleyano, OC₂₋₆alkyleyano, C₀₋₆alkylNR⁵R⁶, OC₂₋₆alkylNR⁵R⁶, C₁₋₆alkyl(CO)NR⁵R⁶, OC₁₋₆alkyl(CO)NR⁵R⁶, C₀₋₆alkylNR⁵(CO)R⁶, OC₂₋₆alkylNR⁵(CO)R⁶, C₀₋₆alkylNR⁵(CO)NR⁵R⁶, C₀₋₆alkylSR⁵, OC₂₋₆alkylSR⁵, C₀₋₆alkyl(SO)R⁵, OC₂₋₆alkyl(SO)R⁵, C₀₋₆alkylSO₂R⁵, OC₂₋₆alkylSO₂R⁵, C₀₋₆alkyl(SO₂)NR⁵R⁶, OC₂₋₆alkyl(SO₂)NR⁵R⁶, C₀₋₆alkylNR⁵(SO₂)R⁶, OC₂₋₆alkylNR⁵(SO₂)R⁶, C₀₋₆alkylNR⁵(SO₂)NR⁵R⁶, OC₂₋₆alkylNR⁵(SO₂)NR⁵R⁶, (CO)NR⁵R⁶, O(CO)NR⁵R⁶, NR⁵OR⁶, C₀₋

~~C₆alkylNR⁵(CO)OR⁶, OC₂₋₆alkylNR⁵(CO)OR⁶, SO₂R⁵ and a 5- or 6-membered ring containing atoms independently selected from the group consisting of C, N, O and S;~~

X¹ is selected from the group consisting ~~[[of:]]~~ of N, NR⁴ and CR⁴;

X² is selected from the group consisting ~~[[of:]]~~ of C and N;

X³ is selected from the group consisting ~~[[of: CR⁴,]]~~ of N and O;

X⁴ is selected from the group consisting ~~[[of:]]~~ of CR⁴, N, NR⁴ and O;

X⁵ is selected from the group consisting ~~[[of:]]~~ of a bond, CR⁴R^{4'}, NR⁴, O, S, SO and SO₂;

X⁶ is selected from the group consisting of: CR⁴ and N;

X⁷ is selected from the group consisting ~~[[of:]]~~ of C and N;

R⁴ and R^{4'} are independently selected from the group consisting of hydrogen, halo, C₁₋₆alkyl and C₁₋₆alkylhalo;

Q is triazolyl;

~~R⁴ is independently selected from a group consisting of hydrogen, hydroxy, C₁₋₆alkyl, C₀₋₆alkylcyano, oxo, =NR⁵, =NOR⁵, C₁₋₄alkylhalo, halo, C₃₋₇cycloalkyl, O(CO)C₁₋₄alkyl, C₁₋₄alkyl(SO)C₀₋₄alkyl, C₁₋₄alkyl(SO₂)C₀₋₄alkyl, (SO)C₀₋₄alkyl, (SO₂)C₀₋₄alkyl, OC₁₋₄alkyl, C₁₋₄alkylOR⁵ and C₀₋₄alkylNR⁵R⁶;~~

~~Q is selected the group consisting of heterocycloalkyl and heteroaryl;~~

R² and R³ are independently selected from the group consisting ~~[[of:]]~~ of hydroxy, C₀₋₆alkylcyano, oxo, =NR⁵, =NOR⁵, C₁₋₄alkylhalo, halo, C₁₋₆alkyl, C₃₋₆cycloalkyl, C₀₋₆alkylaryl, C₀₋₆alkylheteroaryl, C₁₋₆alkylcycloalkyl, C₀₋₆alkylheterocycloalkyl, OC₁₋₄alkyl, OC₀₋₆alkylaryl, O(CO)C₁₋₄alkyl, (CO)OC₁₋₄alkyl, C₀₋₄alkyl(S)C₀₋₄alkyl, C₁₋₄alkyl(SO)C₀₋₄alkyl, C₁₋₄alkyl(SO₂)C₀₋₄alkyl, (SO)C₀₋₄alkyl, (SO₂)C₀₋₄alkyl, C₁₋₄alkylOR⁵, C₀₋₄alkylNR⁵R⁶ and a 5- or 6-membered ring containing atoms independently selected from C, N, O and S, which ring may

optionally be fused with a 5- or 6-membered ring containing atoms independently selected from the group consisting of C, N and O and wherein said ring and said fused ring may be substituted by one or more A;

wherein any C₁₋₆alkyl, aryl, or heteroaryl defined under R¹, R² and R³ may be substituted by one or more A;

A is selected from the group consisting of: hydrogen, hydroxy, halo, nitro, oxo, C₀₋₆alkylcyano, C₀₋₄alkylC₃₋₆cycloalkyl, C₁₋₆alkyl, -OC₁₋₆alkyl, C₁₋₆alkylhalo, OC₁₋₆alkylhalo, C₂₋₆alkenyl, C₀₋₃alkylaryl, C₀₋₆alkylOR⁵, OC₂₋₆alkylOR⁵, C₀₋₆alkylSR⁵, OC₂₋₆alkylSR⁵, (CO)R⁵, O(CO)R⁵, OC₂₋₆alkylcyano, OC₁₋₆alkylCO₂R⁵, O(CO)OR⁵, OC₁₋₆alkyl(CO)R⁵, C₁₋₆alkyl(CO)R⁵, NR⁵OR⁶, C₀₋₆NR⁵R⁶, OC₂₋₆alkylNR⁵R⁶, C₀₋₆alkyl(CO)NR⁵R⁶, OC₁₋₆alkyl(CO)NR⁵R⁶, OC₂₋₆alkylNR⁵(CO)R⁶, C₀₋₆alkylNR⁵(CO)R⁶, C₀₋₆alkylNR⁵(CO)NR⁵R⁶, O(CO)NR⁵R⁶, C₀₋₆alkyl(SO₂)NR⁵R⁶, OC₂₋₆alkyl(SO₂)NR⁵R⁶, C₀₋₆alkylNR⁵(SO₂)R⁶, OC₂₋₆alkylNR⁵(SO₂)R⁶, SO₃R⁵, C₁₋₆alkylNR⁵(SO₂)NR⁵R⁶, OC₂₋₆alkyl(SO₂)R⁵, C₀₋₆alkyl(SO₂)R⁵, C₀₋₆alkyl(SO)R⁵, OC₂₋₆alkyl(SO)R⁵ and a 5- or 6-membered ring containing atoms independently selected from the group consisting of C, N, O and S;

R⁵ and R⁶ are independently selected from, H, C₁₋₆alkyl, C₃₋₇cycloalkyl and aryl;

m is selected from 0, 1, 2, 3 or 4 1 or 2;

n is selected from 0, 1, 2, 3 or 4;

p is selected from 0, 1, 2, 3 or 4; ~~[[and]]~~ or

~~a salt or hydrate thereof;~~

~~with the proviso that the compound is not:~~

~~4,4'-(1,2-piperazinediyl)di-antipyrine;~~

~~4,4'-(1,2-piperazinediyl)di-antipyrine dihydrochloride; or~~

~~4,4'-(1,2-piperazinediyl)di-antipyrine dipicrate;~~

a pharmaceutically acceptable salt thereof.

2. (Canceled)

3. (Original) A compound according to claim 1 wherein X^7 is C.

4. (Withdrawn) A compound according to claim 1 wherein X^5 is selected from the group consisting of $CR^4R^{4'}$, NR^4 , O, S, SO and SO_2 .

5-9. (Canceled)

10. (currently amended) A compound according to claim [[9]] 1 wherein R^1 is selected from the group consisting of: Cl, F, Me, OMe, CF_3 , OCF_3 , and CN.

11. (Original) A compound according to claim 1 wherein X^2 is C.

12. (Original) A compound according to claim 11 wherein X^1 is N or CR^4 .

13. (Original) A compound according to claim 12 wherein when X^3 is O, X^4 is N and when X^3 is N, X^4 is O.

14. (Original) A compound according to claim 1 wherein X^2 is N.

15. **(Original)** A compound according to claim 14 wherein X^1 is N.

16. **(Original)** A compound according to claim 15 wherein X^3 is N and X^4 is N or CR^4 .

17. **(Canceled)**

18. **(Original)** A compound according to claim 12 wherein X^5 is selected from the group consisting of a bond, $CR^4R^{4'}$, NR^4 and O.

19. **(Original)** A compound according to claim 13 wherein X^5 is selected from the group consisting of a bond, O and NR^4 .

20. **(Withdrawn)** A compound according to claim 16 wherein X^5 is selected from the group consisting of O and CR^4 .

21-24. **(Canceled)**

25. **(Currently amended)** A compound according to claim 1 wherein R^2 and R^3 are independently selected from the group consisting ~~[[of:]]~~ of C_{1-4} alkylhalo, C_{1-6} alkyl, C_{3-6} cycloalkyl, C_{0-6} alkylaryl and C_{0-6} alkylheteroaryl.

26. **(Currently amended)** A compound according to claim 1 wherein A is selected from the group consisting of hydrogen, hydroxyl, halo, C₀₋₆alkylcyano, C₁₋₆alkyl, -OC₁₋₆alkyl, C₁₋₆alkylhalo, OC₁₋₆alkylhalo.

27. **(Currently Amended)** A compound according to claim 1 selected from:

4-(5-{2-[5-(3-Chloro-phenyl)-isoxazol-3-yl]-piperidin-1-yl}-4-methyl-4H-[1,2,4]triazol-3-yl)-pyridine,

3-[5-(3-Chloro-phenyl)-isoxazol-3-yl]-4-(4-methyl-5-pyridin-4-yl-4H-[1,2,4]triazol-3-yl)-morpholine,

3-[5-(3-Chloro-phenyl)-isoxazol-3-yl]-4-[5-(4-difluoromethoxy-phenyl)-4-methyl-4H-[1,2,4]triazol-3-yl]-morpholine,

3-[3-(3-Chloro-phenyl)-[1,2,4]oxadiazol-5-yl]-4-(4-methyl-5-pyridin-4-yl-4H-[1,2,4]triazol-3-yl)-morpholine,

3-[3-(3-Chloro-phenyl)-[1,2,4]oxadiazol-5-yl]-4-[5-(4-difluoromethoxy-phenyl)-4-methyl-4H-[1,2,4]triazol-3-yl]-morpholine,

3-[3-(3-Chloro-phenyl)-[1,2,4]oxadiazol-5-yl]-4-(4-methyl-5-pyridin-4-yl-4H-[1,2,4]triazol-3-yl)-piperazine-1-carboxylic acid tert-butyl ester,

2-[3-(3-Chloro-phenyl)-[1,2,4]oxadiazol-5-yl]-1-(4-methyl-5-pyridin-4-yl-4H-[1,2,4]triazol-3-yl)-piperazine,

2-[3-(3-Chloro-phenyl)-[1,2,4]oxadiazol-5-yl]-4-methyl-1-(4-methyl-5-pyridin-4-yl-4H-[1,2,4]triazol-3-yl)-piperazine,

3-[3-(3-Chloro-phenyl)-[1,2,4]oxadiazol-5-yl]-4-[5-(4-difluoromethoxy-phenyl)-4-methyl-4H-[1,2,4]triazol-3-yl]-piperazine-1-carboxylic acid tert-butyl ester,

2-[3-(3-Chloro-phenyl)-[1,2,4]oxadiazol-5-yl]-1-[5-(4-difluoromethoxy-phenyl)-4-methyl-4H-[1,2,4]triazol-3-yl]-piperazine,

2-[3-(3-Chloro-phenyl)-[1,2,4]oxadiazol-5-yl]-1-[5-(4-difluoromethoxy-phenyl)-4-methyl-4H-[1,2,4]triazol-3-yl]-4-methyl-piperazine,

2-[2-(3-Chlorophenyl)-2H-tetrazol-5-yl]-1-{5-[4-(difluoromethoxy)phenyl]-4-methyl-4H-1,2,4-triazol-3-yl}piperidine,

4-(5-{2-[2-(3-chlorophenyl)-2H-tetrazol-5-yl]piperidin-1-yl}-4-methyl-4H-1,2,4-triazol-3-yl)pyridine,

2-[2-(3-Chlorophenyl)-2H-tetrazol-5-yl]-1-[5-(4-methoxyphenyl)-4-methyl-4H-1,2,4-triazol-3-yl]piperidine,

[4-(5-{2-[2-(3-chlorophenyl)-2H-tetrazol-5-yl]piperidin-1-yl}-4-methyl-4H-1,2,4-triazol-3-yl)phenyl]dimethylamine,

[4-(5-{2-[2-(3-Chloro-phenyl)-2H-tetrazol-5-yl]-piperidin-1-yl}-4-methyl-4H-[1,2,4]triazol-3-yl)-benzyl]-dimethyl-amine,

{2-[4-(5-{2-[2-(3-Chloro-phenyl)-2H-tetrazol-5-yl]-piperidin-1-yl}-4-methyl-4H-[1,2,4]triazol-3-yl)-phenoxy]-ethyl}-dimethyl-amine,

(R)-3-[3-(3-Chloro-phenyl)-[1,2,4]oxadiazol-5-yl]-4-(4-methyl-5-pyridin-4-yl-4H-[1,2,4]triazol-3-yl)-morpholine,

(S) 3-[3-(3-Chloro-phenyl)-[1,2,4]oxadiazol-5-yl]-4-(4-methyl-5-pyridin-4-yl-4H-[1,2,4]triazol-3-yl)-morpholine,

(R)-2-[2-(3-Chlorophenyl)-2H-tetrazol-5-yl]-1-{5-[4-(difluoromethoxy)phenyl]-4-methyl-4H-1,2,4-triazol-3-yl}piperidine,

(S)-2-[2-(3-Chlorophenyl)-2H-tetrazol-5-yl]-1-{5-[4-(difluoromethoxy)phenyl]-4-methyl-4H-1,2,4-triazol-3-yl}piperidine,

(R)-4-(5-{2-[2-(3-Chlorophenyl)-2H-tetrazol-5-yl]piperidin-1-yl}-4-methyl-4H-1,2,4-triazol-3-yl)pyridine,

(S)-4-(5-{2-[2-(3-Chlorophenyl)-2H-tetrazol-5-yl]piperidin-1-yl}-4-methyl-4H-1,2,4-triazol-3-yl)pyridine,

4-[5-(5-{2-[5-(3-Chloro-phenyl)-isoxazol-3-yl]-pyrrolidin-1-yl}-4-cyclopropyl-4H-[1,2,4]triazol-3-yl)-pyridin-2-yl]-morpholine,

4-[5-(5-{2-[5-(3-Chloro-phenyl)-isoxazol-3-yl]-pyrrolidin-1-yl}-4-methyl-4H-[1,2,4]triazol-3-yl)-pyridin-2-yl]-morpholine,

3-(5-{2-[5-(3-Chloro-phenyl)-isoxazol-3-yl]-pyrrolidin-1-yl}-4-methyl-4H-[1,2,4]triazol-3-yl)-pyridine,

4-(5-{2-[5-(3-Chloro-phenyl)-isoxazol-3-yl]-pyrrolidin-1-yl}-4-cyclopropyl-4H-[1,2,4]triazol-3-yl)-pyridine,

3-[5-(3-Chloro-phenyl)-[1,2,4]oxadiazol-3-yl]-4-(5-pyridin-4-yl-4H-[1,2,4]triazol-3-yl)-morpholine,

3-[5-(3-chlorophenyl)isoxazol-3-yl]-4-(4-cyclopropyl-5-pyridin-3-yl-4H-1,2,4-triazol-3-yl)morpholine,

3-[5-(3-chlorophenyl)isoxazol-3-yl]-4-(4-cyclopropyl-5-pyridin-4-yl-4H-1,2,4-triazol-3-yl)morpholine,

3-[5-(3-chlorophenyl)isoxazol-3-yl]-4-(4-methyl-5-pyridin-3-yl-4H-1,2,4-triazol-3-yl)morpholine,

3-[5-(3-Chloro-phenyl)-isoxazol-3-yl]-4-[5-(6-methoxy-pyridin-3-yl)-4-methyl-4H-[1,2,4]triazol-3-yl]-morpholine,

3-[3-(3-chlorophenyl)-1,2,4-oxadiazol-5-yl]-4-[5-(2-methoxypyridin-4-yl)-4-methyl-4H-1,2,4-triazol-3-yl]morpholine,

3-[3-(3-chlorophenyl)-1,2,4-oxadiazol-5-yl]-4-[5-(2-methylpyridin-4-yl)-4-methyl-4H-1,2,4-triazol-3-yl]morpholine,

3-[3-(3-chlorophenyl)-1,2,4-oxadiazol-5-yl]-4-[5-(5-fluoropyridin-3-yl)-4-methyl-4H-1,2,4-triazol-3-yl]morpholine,

3-[5-(3-chlorophenyl)isoxazol-3-yl]-4-[5-(5-fluoropyridin-3-yl)-4-methyl-4H-1,2,4-triazol-3-yl]morpholine,

3-[3-(3-chlorophenyl)-1,2,4-oxadiazol-5-yl]-4-(4-methyl-5-pyridin-2-yl-4H-1,2,4-triazol-3-yl)morpholine,

4-[5-(5-fluoropyridin-3-yl)-4-methyl-4H-1,2,4-triazol-3-yl]-3-[3-(3-iodophenyl)-1,2,4-oxadiazol-5-yl]morpholine,

3-[3-(3-iodophenyl)-1,2,4-oxadiazol-5-yl]-4-(4-methyl-5-pyridin-4-yl-4H-1,2,4-triazol-3-yl)morpholine,

3-[5-(3-chlorophenyl)isoxazol-3-yl]-4-[5-(2-methylpyridin-4-yl)-4-methyl-4H-1,2,4-triazol-3-yl]morpholine,

3-[2-(3-chlorophenyl)-2H-tetrazol-5-yl]-4-(4-methyl-5-pyridin-3-yl-4H-1,2,4-triazol-3-yl)morpholine,

3-[2-(3-chlorophenyl)-2H-tetrazol-5-yl]-4-[5-(3,5-difluorophenyl)-4-methyl-4H-1,2,4-triazol-3-yl]morpholine,

3-(5-{2-[5-(3-chlorophenyl)isoxazol-3-yl]pyrrolidin-1-yl}-4-cyclopropyl-4H-1,2,4-triazol-3-yl)pyridine, and

4-(5-{2-[5-(3-chlorophenyl)isoxazol-3-yl]pyrrolidin-1-yl}-4-methyl-4H-1,2,4-triazol-3-yl)pyridine.

28. (Currently amended) A pharmaceutical composition comprising as active ingredient a therapeutically effective amount of the compound according to any one of claims [[1

to 26,]] 1, 3, 4, 10-16, 18-20, 25 and 26, in association with one or more pharmaceutically acceptable diluent, excipients and/or inert carrier.

29. (Canceled)

30. (Previously Presented) The compound according to claim 1, for use in therapy.

31. (Previously Presented) The compound according to claim 1, for use in treatment of mGluR 5 mediated disorders.

32. (Withdrawn) Use of the compound according to claim 1, in the manufacture of a medicament for the treatment of mGluR 5 mediated disorders.

33. (Currently Amended - Withdrawn) A method of treatment of mGluR 5 mediated disorders, comprising ~~administering~~ administering to a mammal, including man in need of such treatment, a therapeutically effective amount of the compound according to claim 1.

34. (Currently Amended - Withdrawn) The method according to claim 33, ~~for use in treatment of~~ wherein the disorders mediated by mGluR 5 are neurological disorders.

35. (Currently Amended - Withdrawn) The method according to claim 33, ~~for use in treatment of~~ wherein the disorders mediated by mGluR 5 are psychiatric disorders.

36. **(Currently Amended - Withdrawn)** The method according to claim 33, ~~for use in treatment of~~ wherein the disorders mediated by mGluR 5 are chronic and acute pain disorders.

37. **(Currently Amended - Withdrawn)** The method according to claim 33, ~~for use in treatment of~~ wherein the disorders mediated by mGluR 5 are gastrointestinal disorders.

38. **(Withdrawn)** A method for inhibiting activation of mGluR 5 receptors, comprising treating a cell containing said receptor with an effective amount of the compound according to claim 1.